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Annli	Application Data Sheet 37 CFF			CED	Attorney Docket Number				101367-1P US					
Appii		Jala S	illeet 37		1.70	Application	on Numb	er						
Title of	f Invention	THI	OPHENE I	DERIVA	ATIVES	AS CHK 1 IN	NHIBITOF	RS						
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	Susan								Ashwe	ell				
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Δnnli	Application Data Sheet 37 CF			CER	Attorney Docket Number				101367-1P US				
Appli	cation Da	ta Sile	JC 1 37	CI IX	1.70	Applica	ation N	lumbe	r				
Title of	Invention	THIOP	HENE C	ERIVA	TIVES	AS CHK 1	INHIB	BITORS	6				
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Applic	ant 4					-						Remove	
	ant Authori	tv •In	ventor		gal Rep	resentativ	e unde	er 35 U	J.S.C. 11	7 (Party of In	terest under 35 U.S.	.C. 118
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Applic	ant 5	•										Remove	
Applic	ant Authori	ty ⊙ln\	ventor	○Le	gal Rep	resentativ	e unde	er 35 U	J.S.C. 11	7 (Party of In	terest under 35 U.S.	.C. 118
Prefix	Given Nan				Middle Name					Famil	Suffix		
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Annli	Application Data Sheet 37 CFR 1.70					Attorney Docket Number			101367-1P US				
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Title of	Invent	ion THI	OPHENE [DERIVAT	IVES /	AS CHK 1	INHIE	BITORS	5				
Mailing	g Addr	ess of Ap	plicant:										
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Applic	ant Au	thority 🖲	Inventor	○Leg	al Rep	Representative under 35 U.S.C. 11			Party of Interest under 35 U.s			C. 118	
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Application Data Sheet 37 CFR 1.76			Attorn	ey Doo	cket N	umber	101367	101367-1P US					
Application Da	ala S	meet 37 CFK 1.76	Applic	ation N	Numbe	er							
Title of Invention	THIC	OPHENE DERIVATIVES	AS CHK 1 INHIBITORS										
Mailing Address	of Apı	olicant:											
Address 1			straZeneca R & D Boston										
Address 2		35 Gatehouse Drive											
City Waltha	am				Stat	e/Provin	ice	M	A				
Postal Code		02451		Cou	ıntryi	US							
	Listed - Additional In m by selecting the Add		Inform	ation	blocks	may be				Add			
Corresponde	orrespondence Information:												
	Enter either Customer Number or complete the Correspondence Information section below. For further information see 37 CFR 1.33(a).												
☐ An Address	An Address is being provided for the correspondence Information of this application.												
Customer Number	er	44992											
Email Address		carol.loeschorn@as	trazeneca	a.com					Add	Email		Remov	e Email
Application Information:													
Title of the Inven	tion	THIOPHENE DERIV	/ATIVES	AS CH	K 1 IN	HIBITORS	3						
Attorney Docket	Numb	er 101367-1P US			S	mall Ent	ity Statu	ıs (Claim	ed [
Application Type	!	Nonprovisional			•								
Subject Matter		Utility											
Suggested Class	(if an	y) N/A			S	ub Clas	s (if any) \	I/A				
Suggested Techi	nology	Center (if any)	N/A					•					
Total Number of	Drawi	ng Sheets (if any)			S	uggeste	d Figure	e fo	r Pub	licatio	on (if	any)	N/A
Publication Infor	matio	ո։			•								
Request Earl	y Publ	ication (Fee required a	t time of	Reque	est 37	CFR 1.2	!19)						
Request Not to Publish. I hereby request that the attached application not be published under 35 U.S.C. 122(b) and certify that the invention disclosed in the attached application has not been and will not be the subject of an application filed in another country, or under a multilateral agreement, that requires publication at eighteen months after filing.													
Representative Information:													
this information in the Enter either Cu	Representative information should be provided for all practitioners having a power of attorney in the application. Providing this information in the Application Data Sheet does not constitute a power of attorney in the application (see 37 CFR 1.32). Enter either Customer Number or complete the Representative Name section below. If both sections are completed the Customer Number will be used for the Representative Information during processing.												
Please Select One	e :	Customer Number	r C) USP	atent F	Practitione	er 🔘	U	S Rep	resenta	ative (37 CFF	R 11.9)
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Application Da	ta Sheet 37 CFR 1.76	Attorney Docket Number	101367-1P US
Application Da	ita Sileet 37 Cl IX 1.70	Application Number	
Title of Invention	THIOPHENE DERIVATIVES A	AS CHK 1 INHIBITORS	

Domestic Priority Information:

This section allows for the applicant to claim benefit under 35 U.S.C. 119(e), 120, 121, or 365(c). Providing this information in the application data sheet constitutes the specific reference required by 35 U.S.C. 119(e) or 120, and 37 CFR 1.78(a)(2) or CFR 1.78(a) (4), and need not otherwise be made part of the specification.

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Prior Application Status			Remove						
Application Number	Continuity Type	Prior Application Number	Filing Date (YYYY-MM-DD)						
	a 371 of international	PCT/GB2004/005400	2004-12-24						
Prior Application Status			Remove						
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PCT/GB2004/005400	non provisional of	60/534310	2004-01-05						
Prior Application Status			Remove						
Application Number	Continuity Type	Prior Application Number	Filing Date (YYYY-MM-DD)						
PCT/GB2004/005400	non provisional of	60/553305	2004-03-15						
Additional Domestic Priority Data may be generated within this form by selecting the Add button.									

Foreign Priority Information:

This section allows for the applicant to claim benefit of foreign priority and to identify any prior foreign application for which priority is not claimed. Providing this information in the application data sheet constitutes the claim for priority as required by 35 U.S.C. 119(b) and 37 CER 1 55(a)

alid 57 CFR 1.35(a).									
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Application Number	Country i	Parent Filing Date (YYYY-MM-DD)	Priority Claimed						
			Yes No						
Additional Foreign Priority Data may be generated within this form by selecting the Add button.									

Assignee Information:

Providing this information in the application data sheet does not substitute for compliance with any requirement of part 3 of Title 37 of the CFR to have an assignment recorded in the Office. Remove Assignee 1 If the Assignee is an Organization check here. $|\mathbf{x}|$

Organization Name AstraZeneca AB										
Mailing Address Information:										
Address 1										
Address 2	SE-151 85									
City	Sodertalje	State/Province								
Country i SE		Postal Code								
Phone Number	46 8 553 260 00	Fax Number	46 8 553 288 20							
Fmail Address	formalities se@astrazeneca.com	•								

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Application Da	ta Sheet 37 CFR 1.76	Attorney Docket Number	101367-1P US						
Application Da	ita Sileet 37 Cl K 1.70	Application Number							
Title of Invention	THIOPHENE DERIVATIVES	AS CHK 1 INHIBITORS							
Additional Assignee Data may be generated within this form by selecting the Add button.									

Signature:

_	A signature of the applicant or representative is required in accordance with 37 CFR 1.33 and 10.18. Please see 37 CFR 1.4(d) for the form of the signature.											
Signature	/Carol A. Loeschorn/	Date (YYYY-MM-DD)	2006-11-30									
First Name	Carol	Registration Number	35590									

This collection of information is required by 37 CFR 1.76. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 23 minutes to complete, including gathering, preparing, and submitting the completed application data sheet form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. **SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.**

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patent application or patent. Accordingly, pursuant to the requirements of the Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) furnishing of the information solicited is voluntary; and (3) the principal purpose for which the information is used by the U.S. Patent and Trademark Office is to process and/or examine your submission related to a patent application or patent. If you do not furnish the requested information, the U.S. Patent and Trademark Office may not be able to process and/or examine your submission, which may result in termination of proceedings or abandonment of the application or expiration of the patent.

The information provided by you in this form will be subject to the following routine uses:

- 1. The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these records.
- 2. A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiations.
- 3. A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
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- 5. A record related to an International Application filed under the Patent Cooperation Treaty in this system of records may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant to the Patent Cooperation Treaty.
- 6. A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or his/her designee, during an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2904 and 2906. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- 8. A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S.C. 122(b) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspections or an issued patent.
- A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.